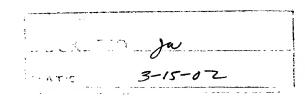
PATENT COOPERATION TREATY

	`			
•		4 r	NES	
PA	ATENT COOPE	RATION TR	EATY	
From the INTERNATIONAL PRELIMINARY EX.	AMINING AUTHORITY		Response to	
To: NICOLAS E. SECKEL E C ARMSTRONG, WESTERMAN, HATT	EIVED		PCT Due: May 12	,'02
MCLELAND & NAUGHTON, LLP 1725 K STREET NW MAR WASHINGTON, DC 20006 MAR	1 5 3002	/	WRITTEN OPINION	
ARMSTRONG,	WESTERMAN, HATTOFI	1,	(PCT Rule 66)	
	LLP	Date of Mailing (day/month/year)	12 MAR 2002	
Applicant's or agent's file reference		REPLY DUE	within 2 months/days from	•
000152PCT International application No.	International filing date		the above date of mailing Priority date (day/month/year)	
PCT/US01/09590	26 March 2001 (26.03.2		24 March 2090 (24: 73.2000)	
International Patent Classification (IPC)	•		24 Par Sil 1030 (25 1.2000)	
IPC(7): A61K 48/00, 38/00, 35/00; C12	N 15/63 and US Cl.: 514/	/2, 44;424/93.21;435	5/320.1	
Applicant				
TKACHUK, ZENOVIY				
 This written opinion is the <u>fir</u> This opinion contains indicati 			eliminary Examining Authority.	
- N-7	_	ng tems.	•	
I Basis of the opinion	on		,	
II Priority				
III Non-establishmen	t of opinion with regard to	o novelty, inventive s	step and industrial applicability	
IV Lack of unity of i	nvention		-	
	ent under Rule 66.2 (a)(ii) anations supporting such s	-	lty, inventive step or industrial applicability;	
VI Certain document	s cited			1
VII Certain defects in	the international application	ion		
VIII Certain observation	ons on the international app	plication		
3. The applicant is hereby invit				
	limit indicated above. The y to grant an extension. Se		ore the expiration of that time limit, request	
•	g a written reply, accompa and the language of the ar		riate, by amendments, according to Rule 66.3. es 66.8 and 66.9.	
For the exam	onal opportunity to submit niner's obligation to considual mal communication with the	der amendments and/	or arguments, see Rule 66.4 bis.	
If no reply is filed, the inter	national preliminary exam	nination report will b	be established on the basis of this opinion.	
4. The final date by which the i examination report must be e		ule 69 .2 is: 24 July ?	2002 (24.07.2002)	
Name and mailing address of the IPEA Commissioner of Patents and Trademar Box PCT		Authorized office	e Bridges for	
Washington, D.C. 20231 Facsimile No. (703)305-3230		Telephone No. 7	703-308-0196	

Form PCT/IPEA/408 (cover sheet)(July 1998)



WRITTEN OPINION

International application No.

PCT/US01/09590

I.	Basis of the opinion	
1.	With regard to the elements of the international application:*	
	the international application as originally filed	
	the description:	
	pages 1-58 , as originally filed	
	pages NONE , filed with the demand	
	pages NONE, filed with the letter of	
	the claims:	
	pages 59-62 , as originally filed	
	pages NONE , as amended (together with any statement) under Article 19	
	pages NONE , filed with the demand	
	pages NONE , filed with the letter of	
	the drawings:	
	pages none, as originally filed	
	pages NONE , filed with the demand	
	pages NONE , filed with the letter of	
	the sequence listing part of the description:	
	pages NONE , as originally filed	
	pages NONE , filed with the demand	
	pages NONE , filed with the letter of	
2.	With regard to the language, all the elements marked above were available or furnished to this Au language in which the international application was filed, unless otherwise indicated under this item. These elements were available or furnished to this Authority in the following language	n.
	the language of a translation furnished for the purposes of international search (under Rule23	.1(b)).
	the language of publication of the international application (under Rule 48.3(b)).	
	the language of the translation furnished for the purposes of international preliminary examin 55.2 and/or 55.3).	ation(under Rules
3.	With regard to any nucleotide and/or amino acid sequence disclosed in the international applicat opinion was drawn on the basis of the sequence listing:	ion, the written
	contained in the international application in printed form.	
	filed together with the international application in computer readable form.	
	furnished subsequently to this Authority in written form.	
	furnished subsequently to this Authority in computer readable form.	
	The statement that the subsequently furnished written sequence listing does not go beyond th	e disclosure in the
	international application as filed has been furnished.	•
	The statement that the information recorded in computer readable form is identical to the writing been furnished.	tten sequence listing
4.	. The amendments have resulted in the cancellation of:	
	the description, pages NONE	
	the claims, Nos. NONE	
	the drawings, sheets/fig NONE	
5.		considered to go
٥.	beyond the disclosure as filed, as indicated in the Supplemental Box (Rule 70.2(c)).	<i>O</i>
	Replacement sheets which have been furnished to the receiving Office in response to an invitation under Articular is opinion as "originally filed."	e 14 are referred to in

WRITTEN OPINION

International application No.

PCT/US01/09590

Ш	. No	on-establishment of opinion with regard to novelty, inventive step and industrial applicability
		question whether the claimed invention appears to be novel, to involve an inventive step (to be non-obvious), or industrially applicable have not been examined in respect of:
		the entire international application,
	\boxtimes	claims Nos. <u>14-21</u>
		because:
		the said international application, or the said claim Nos relate to the following subject matter which does not require international preliminary examination (specify):
		•
	\boxtimes	the description, claims or drawings (indicate particular elements below) or said claims Nos. 14-21 are so unclear that no meaningful opinion could be formed (specify):
bec	ause t	they are dependent claims and are not drafted in accordance with the second and third sentences of Rule.
		the claims, or said claims Nos are so inadequately supported by the description that no meaningful opinion could be formed.
	\boxtimes	no international search report has been established for said claims Nos. 14-21.
		ritten opinion cannot be drawn due to the failure of the nucleotide and/or amino acid sequence listing to comply the standard provided for in Annex C of the Administrative Instructions:
		the written form has not been furnished or does not comply with the standard.
		the computer readable form has not been furnished or does not comply with the standard.

Form PCT/IPEA/408 (Box III) (July 1998)

WRITTEN OPINION

International application No. PCT/US01/09590

v.	Reasoned statement under Rule 66.2(a)(ii) with regard to novelty, inventive step or industrial applicability;
	citations and explanations supporting such statement

1. STATEMENT

Novelty (N)	Claims	30-33	YES
•	Claims	1-13, 22-29	NO
Inventive Step (IS)	Claims	NONE	YES
-	Claims	1-13, 22-33	NO
Industrial Applicability (IA)	Claims	1-13, 22-33	YES
• • •	Claims	NONE	NO

2. CITATIONS AND EXPLANATIONS

Claims 1-11, and 22-29 lack novelty under PCT Article 33(2) as being anticipated by WO 94/02595.

These claims are directed to method for the treatment of inflammation or inflammatory-related disorder comprising administering to a mammal in need of an effective amount of ribonucleic acid and a pharmaceutically acceptable vehicle, wherein the inflammatory-associated disorder is infarct, stroke, arthritis, allergy, etc., and the composition used in the method.

WO 94/02595 teaches using an enzymatic RNA molecule for treating inflammatory associated disorders, such as arthritis, psoriasis, and cardiovascular diseases (page 10 and claims 1, 2, 11, and 12). Thus, WO 94/02595 anticipates these claims.

Claims 1-13, 26, and 29 lack novelty under PCT Article 33(2) as being anticipated by US 5,712,256.

These claims are directed to method for the treatment of inflammation or inflammatory-related disorder comprising administering to a mammal in need of an effective amount of ribonucleic acid and a pharmaceutically acceptable vehicle, wherein the inflammatory-associated disorder is pain, and swelling, etc., and the composition used in the method, wherein the dosing ranges from 0.1mg to 1g/kg, or 0.1 to 1 gram.

US 5,712,256 teaches using an enzymatic RNA molecule for treating inflammatory associated disorders, such as wound (claims 1, 2, 3), wherein the sufficient dose is about 0.17g/kg per day (claim 4). Thus, WO 94/02595 anticipates these claims.

Claims 1-13, 22-33 lack an inventive step under PCT Article 33(3) as being obvious over the prior art as applied in the immediately preceding paragraph and further in view of US 3,615,654.

Claims 31-33 further drawn to the composition wherein the ribonucleic acid has a nitrogen content of more than 14.5% by weight, and a phosphorus content of more than 8.5% by weight, wherein the pharmaceutical carrier is capsules or suppositories.

US 3,615,654 teaches that pure ribonucleic acids from microbial cells have 8.5% phosphorus and 15.1% nitrogen (column 7, lines 24-25). And the specification teaches to obtain the RNA from microbial cells. Further capsules and suppositories are the commonly used pharmaceutical carriers. Thus, claimed invention, as a whole is obvious over the cited prior art.

Claims 1-13, 22-33 meet the criteria set out for industrial applicability in PCT Article 33(4), because the method could be used treating inflammatory associated disorders.

	NEW CITATIONS
US 3 615 654 A /	(AVIJKAWA V) 26 October 1971, column 7, lines 24-25). Relevant to claims 31 and 32.

WR	ITTEN	OPIN	JION

International application No. PCT/US01/09590

TIME LIMIT: The time limit set for response to a Written Opinion may not be extended. 37 CFR 1.484(d). Any response received after the expiration of the time limit set in the Written Opinion will not be considered in preparing the International Preliminary Examination Report.

Form PCT/IPEA/408 (Supplemental Box) (July 1998)